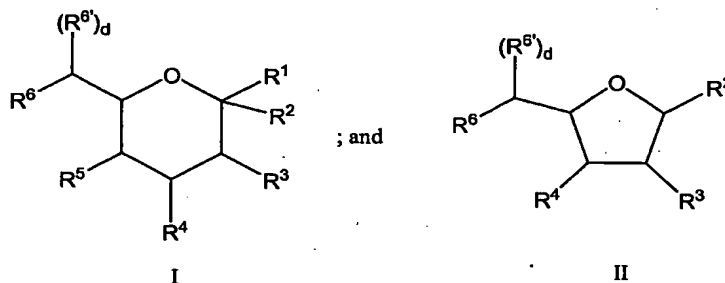


**WHAT IS CLAIMED IS:**

- 1    1.    A compound having a formula that is a member selected from:



3 wherein

4  $R^1$  is H,  $CH_2OR^7$ ,  $COOR^7$  or  $OR^7$

5 in which

6 R<sup>7</sup> represents H, substituted or unsubstituted alkyl or substituted or  
7 unsubstituted heteroalkyl;

8 R<sup>2</sup> is a member selected from H, OH, an activating group and a moiety that includes a  
9 nucleotide;

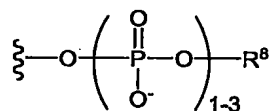
10  $R^3, R^4, R^5, R^6$  and  $R^{6'}$  are independently selected from H, substituted or unsubstituted  
11 alkyl,  $OR^9$ , and  $NHC(O)R^{10}$

12                    wherein

13 R<sup>9</sup> and R<sup>10</sup> are independently selected from H, substituted or unsubstituted  
14 alkyl or substituted or unsubstituted heteroalkyl,

15 and at least one of  $R^3$ ,  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^{6'}$  includes a polymeric modifying moiety.

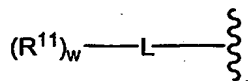
- 1      2.      The compound according to claim 1 wherein R<sup>2</sup> has the formula:



3 in which R<sup>8</sup> is a nucleoside.

- 1     3.     The compound according to claim 2 wherein R<sup>8</sup> is a member selected from cytosine,  
2     uridine, guanosine, adenosine and thymidine.

4. The compound according to claim 1 wherein at least one of  $R^3$ ,  $R^4$ ,  $R^5$  and  $R^6$  includes the moiety:



wherein

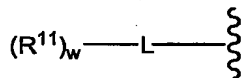
$R^{11}$  is a polymeric modifying moiety;

$L$  is a member selected from a bond and a linking group; and

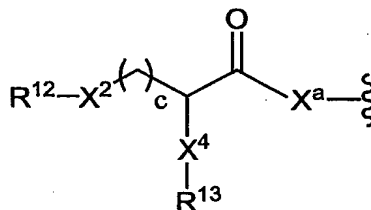
$w$  is selected from the integers from 1 to 6.

5. The compound according to claim 4 wherein said linking group is a member selected from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl moieties.

6. The compound according to claim 5 wherein the moiety:



has the formula:



wherein

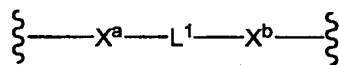
$\text{X}^2$  and  $\text{X}^4$  are independently selected from linkage fragments;

$\text{X}^a$  is a linkage fragment;

$\text{R}^{12}$  and  $\text{R}^{13}$  are independently selected polymeric arms; and

$c$  is an integer from 1 to 20.

7. The compound according to claim 5 wherein said linking group has the formula:



in which

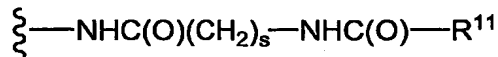
$\text{X}^a$  and  $\text{X}^b$  are independently selected linkage fragments; and

$\text{L}^1$  is a member selected from a bond, substituted or unsubstituted alkyl or substituted or unsubstituted heteroalkyl.

1 8. The compound according to claim 7 wherein  $X^a$  and  $X^b$  are linkage fragments  
 2 independently selected from S, SC(O)NH, HNC(O)S, SC(O)O, O, NH, NHC(O), (O)CNH  
 3 and NHC(O)O, and OC(O)NH.

1 9. The compound according to claim 5 wherein said linker comprises an acyl moiety.

1 10. The compound according to claim 9 wherein  $L-R^{11}$  has the formula:

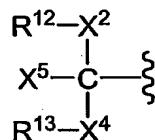


3 in which

4 s is an integer from 0 to 20; and

5  $R^{11}$  is said polymeric modifying moiety.

1 11. The compound according to claim 1, wherein said polymeric modifying moiety has  
 2 the formula:



4 wherein

5  $X^2$  and  $X^4$  are independently selected from linkage fragments;

6  $X^5$  is a non-reactive group; and

7  $R^{12}$  and  $R^{13}$  are independently selected polymeric arms.

1 12. The compound according to claim 11 wherein  $X^2$  and  $X^4$  are linkage fragments  
 2 independently selected from S, SC(O)NH, HNC(O)S, SC(O)O, O, NH, NHC(O), (O)CNH  
 3 and NHC(O)O, OC(O)NH and  $(\text{CH}_2)_g Y''$

4 wherein

5 g is an integer from 1 to 50; and

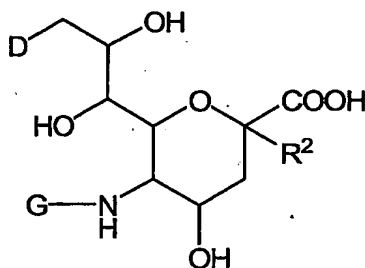
6  $Y''$  is a member selected from O, S and NH.

1 13. The compound according to claim 11 wherein

2  $X^4$  is a peptide bond; and

3  $R^{13}$  is an amino acid residue.

1 14. The compound according to claim 1 having the formula:



in which

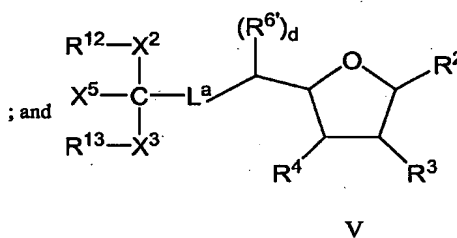
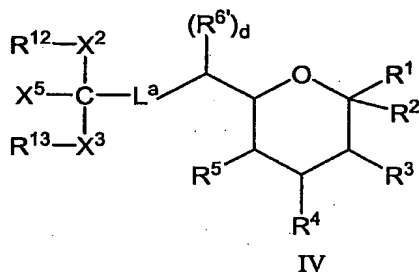
D is a member selected from -OH and  $(R^{11})_{w'}-L-$ ;

G represents is a member selected from H,  $(R^{11})_{w'}-L-$  and  $-C(O)(C_1-C_6)alkyl$ ;

$w'$  is an integer from 2 to 6, and

at least one of D and G is  $(R^{11})_{w'}-L-$ .

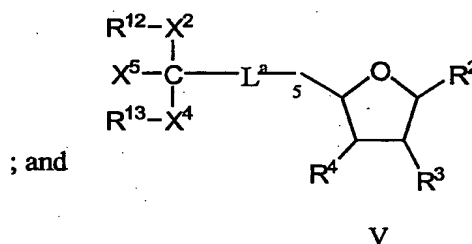
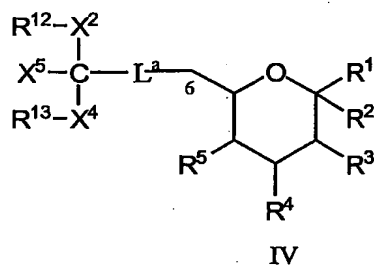
15. The compound according to claim 14 having the formula:



wherein

$L^a$  is a member selected from substituted or unsubstituted alkyl and substituted or unsubstituted heteroalkyl.

16. The compound according to claim 1 having the formula:



wherein

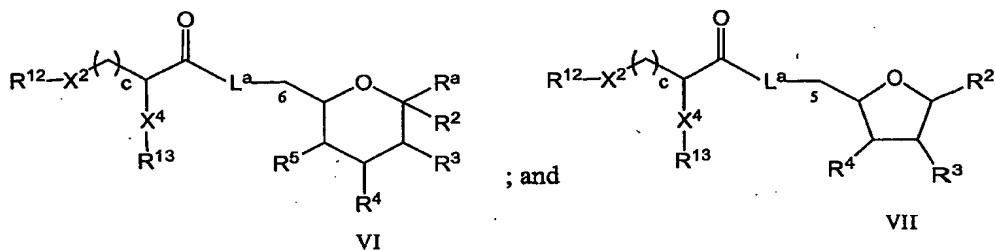
$L^a$  is a member selected from an amino acid residue and a peptidyl residue having from 2 to 4 amino acid residues;

$X^2$  and  $X^4$  are independently selected from linkage fragments;

7  $X^5$  is a non-reactive group; and

8  $R^{12}$  and  $R^{13}$  are independently selected polymeric arms

1    17.    The compound according to claim 16 having the formula:



2  
3 wherein

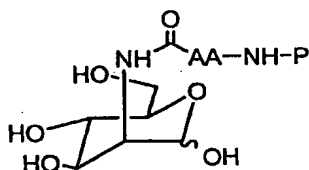
4  $X^2$  and  $X^4$  are independently selected from linkage fragments;

5  $X^a$  is a linkage fragment;

6  $R^{12}$  and  $R^{13}$  are independently selected polymeric arms; and

7       $c$  is an integer from 1 to 20.

1    18.    The compound according to claim 1, having the formula:



2  
3 wherein

4 AA-NH is an amino acid residue; and

5 P is a polymeric modifying group.

1 19. The compound according to claim 18 wherein -AA-NH is -CH<sub>2</sub>NH.

1    20.    The compound according to claim 1 wherein said compound is a substrate for an  
2    enzyme that transfers a sugar moiety from a member selected from an activated sugar, a  
3    nucleotide sugar and combinations thereof onto an acceptor moiety of a substrate.

1    **21.**    The compound according to claim 20 wherein said acceptor moiety is a member  
2    selected from a glycosyl residue, an amino acid residue and an aglycone.

1 22. A method of preparing cytidine monophosphate sialic acid-poly(ethylene glycol), said  
2 method comprising:

3 (a) contacting mannosamine with an activated, N-protected amino acid  
4 under conditions appropriate to form an amide conjugate between said mannosamine and the  
5 N-protected amino acid;

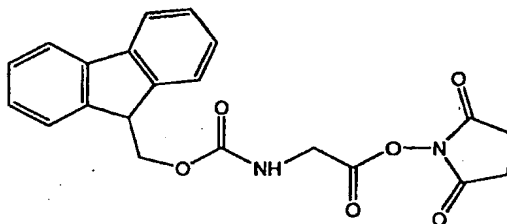
6 (b) contacting said amide conjugate with pyruvate and sialic acid aldolase  
7 under conditions appropriate to convert said amide conjugate to a sialic acid amide conjugate;

8 (c) contacting said sialic acid amide conjugate with cytidine triphosphates,  
9 and a synthetase under conditions appropriate to form a cytidine monophosphate sialic acid  
10 amide conjugate;

11 (d) removing the N-protecting group from said cytidine monophosphate  
12 sialic acid amide conjugate, thereby producing a free amine; and

13 (e) contacting said free amine with an activated PEG, thereby forming said  
14 cytidine monophosphate sialic acid-poly(ethylene glycol).

1 23. The method according to claim 21, wherein said activated N-protected amino acid has  
2 the formula:



3